Integrity test pharma vial

FI21EN14A APPLICATION PHARMA VIAL

**ZERO WASTE** 

NON-DESTRUCTIVE

Application : Pharma vial

Leaks to be detected : Integrity fault (stopper + crimped cap)

**Specifications :** Sample test, deterministic. Reference : ISO 8871-5 (dye test)

Solution : Integrity tester ASC 7400F

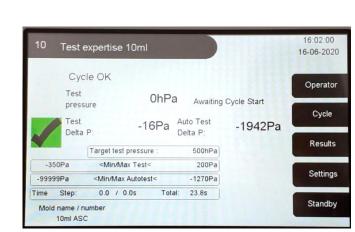
**Operation :** Operator places the vial in the test chamber, brings down the upper part of the chamber and presses « Start ». The instrument locks the test chamber and the cycle starts. The test pressure (positive or vacuum) is established in the chamber. After a short stabilisation, the pressure decay ( $\Delta P$ ) is measured. This  $\Delta P$  is compared to a reject level : a value above this limit indicates a non-conform integrity. The red light comes on and the instrument stays locked. To unlock it, the operator must press an "Unlock process" button.

A  $\Delta P$  below the reject limit means the integrity level is correct. The instrument carries out a certification stage involving an internal calibrated leak, which validates the result instantly : the green light comes on and the chamber is unlocked.

ASC 7400F – Integrity tester for vials/syringes

Cycle time : approx. 10-20 s

Typical reject level : according to standard (ISO 8871-5, ASTM F2338-09, ...) or any "in-house" standard. (Transposition report at request.)



Screen showing a correct integrity result :  $\Delta P = -16 Pa$  for a reject level at -350 Pa







Pharma vial